

**Table S1. Oligonucleotides used in this study**

<b>Primers</b>	<b>Sequences (5'-3')</b>
<b><i>For amplification of CDS</i></b>	
<i>HcSMA2-F</i>	ATGAGCATACAAACGAAG
<i>HcSMA2-R</i>	ATGAGCATACAAACGAAG
<b><i>For the preparation of dsRNA</i></b>	
<i>Hcsma2i-F1</i>	<b>TAATACGACTCACTATAGGGAGACATTGGGCGACAATCAGCTA</b>
<i>Hcsma2i-R1</i>	<u>AAGCTT</u> ATGGATGAGATCGGTTCGAGG
<i>Hcsma2i-F2</i>	<u>AAGCTT</u> CATTGGGCGACAATCAGCTA
<i>Hcsma2i-R2</i>	<b>TAATACGACTCACTATAGGGAGAATGGATGAGATCGGTTCGAGG</b>
<i>Btcry1Ac-F1</i>	<b>TAATACGACTCACTATAGGGCCAATACAGTACCAGCTACAG</b>
<i>Btcry1Ac-R1</i>	<u>GGATCC</u> GATTTCGGCTCTCCACAC
<i>Btcry1Ac-F2</i>	<u>GGATCC</u> CCAATACAGTACCAGCTACAG
<i>Btcry1Ac-R2</i>	<b>AATACGACTCACTATAGGGGATTTCGGCTCTCCACAC</b>
<b><i>For real-time PCR</i></b>	
<i>rtHcsma2-F</i>	ATCCCACCAGGAATTCGACG
<i>rtHcsma2-R</i>	CATCGTCCACGTGCATTTGA
<i>Tubulin-F</i>	TGTTCCATCACCCAAGGTATCC
<i>Tubulin-R</i>	TGACAGACACAAGGTGGTTGAGAT
<i>18S-F</i>	AATGGTTAAGAGGGACAATTCG
<i>18S-R</i>	CTTGGCAAATGCTTTTCGC

**For bimolecular fluorescence complementation plasmids**

<i>Hcdaf8</i> -HA-F	TGAACCGTCAGATCC <b><u>GCTAGCC</u></b> CACCATGCGATCCTTGTTCGAATC
<i>Hcdaf8</i> -HA-R	GCACATCGTAAGGATAT <b><u>CTCGAG</u></b> CCCGTAAATGAGGAATGTGCCA
<i>Hcsma2</i> -Myc-F	TGAACCGTCAGATCC <b><u>GCTAGCC</u></b> CACCATGAGCATACAAACGAAGTT
<i>Hcsma2</i> -Myc-R	TGATCAGCTTCTGCTCG <b><u>CCGATCG</u></b> CTGAAATGGATGAGATCGGTC

Underlined base pairs represent restriction sites, and those in bold letters indicate the T7 promoter site.

**Table S2. Sequences of siRNA used for RNA interference (5'-3')**

siRNAs	Sense strand	Antisense strand	Position of the target sequence
siRNA-136	GCCAGUCUGGAAUUCGCAUTT	AUGCGAAUUCAGACUGGCTT	136-154
siRNA-625	CCAGAACAUUGGGCGACAATT	UUGUCGCCCAAUGUUCUGGTT	625-644
siRNA-1130	GCUGGAUCGAAAUUCAUUUTT	AAAUGAAUUUCGAUCCAGCTT	1130-1149
Negative siRNA	UUCUCCGAACGUGUCACGUTT	ACGUGACACGUUCGGAGAATT	/